

10/502244

## A novel target to inhibit angiogenesis

### Field of the invention

The invention relates to the field of angiogenesis. In particular the invention relates to the use of molecules binding to prominin-1 that can be used for the manufacture of a medicament to prevent pathological angiogenesis.

### Background of the invention

Prominin-1 (PROM-1), also called AC133 or recently designed CD133 (National Center for Biotechnology, 2000), is a rather novel human hematopoietic stem cell antigen <sup>1</sup> of unknown physiological or pathological function. Prominin-1-antigen was first detected on CD34<sup>bright</sup> hematopoietic stem cells <sup>2</sup> and has since been widely used to facilitate the analysis and isolation of hematopoietic and primitive cells <sup>3-5</sup>. Only few prominin-1<sup>+</sup> cells do not coexpress CD34: these cells are very small and define a population of unknown delineation <sup>6</sup>. In acute myeloid leukemias, PROM-1 expression is often but not always associated with CD34 expression <sup>7,8</sup>. Prominin-1 is also found on acute lymphoid leukemia blasts and on a subset of CD34<sup>+</sup> B-cell precursors <sup>9</sup>. Flow cytometry analyses of a wide panel of human cell lines showed that only retinoblastoma and teratocarcinoma cell lines express prominin-1<sup>10</sup>. More recently, it was shown that endothelial progenitor cells co-express PROM-1 antigen and the endothelial cell-specific receptor kinase-inert domain-containing acceptor (KDR) in subpopulations of CD34<sup>+</sup> cells derived from fetal liver, bone marrow, cord blood and peripheral blood <sup>11,12</sup>. Recently, human central nervous system stem cells were also reported to express prominin-1-antigen <sup>13</sup>. A characteristic feature of this protein is its rapid down-regulation during cell differentiation <sup>12,14</sup>, which makes it a unique cell surface marker for the identification and isolation of stem cells and progenitor cells. Human PROM-1 antigen is a glycoprotein of 120 KD and contains an extracellular N- terminus, two extracellular loops, five transmembrane domains, two small cysteine-rich cytoplasmic loops and a cytoplasmic C terminus <sup>1</sup>. Recently a novel isoform of human PROM-1 with a 27 basepair deletion has been described <sup>15</sup>. A structural and sequence-related protein, was identified as the mouse orthologue of human PROM-1 <sup>14</sup>. The 5-transmembrane structure appears phylogenetically conserved from mammals to zebrafish and in fruit flies and nematodes <sup>16,17</sup>. Murine prominin-1, which has a 65% amino acid homology

with human PROM-1 also exists in two isoforms. The short human and murine prominin isoform both encode proteins that lack a 9-amino acid segment at the same location in the N-terminal extracellular region just proximal to the first transmembrane domain<sup>15,18,19</sup>. Although human PROM-1 has been used as a cell surface marker to identify and isolate certain stem cell and progenitor cell populations, the molecular mechanism of how this protein functions remain unclear. The possible role of PROM-1 in hematopoiesis and vasculogenesis in the developing embryo and, after birth, in angiogenesis, postnatal vasculogenesis and hematopoietic stem cell trafficking, remains largely unknown. To study in detail the *in vivo* role of PROM-1 in the present invention PROM-1 deficient mice were generated. It was surprisingly found that PROM-1 has a key role in pathological angiogenesis and that inhibitors of PROM-1 can be used in therapeutic strategies to inhibit blood vessel formation in various pathological disorders.

#### Aims and detailed description of the invention

The "hemangioblast" is a putative progenitor cell that has the potential to form either endothelial or hematopoietic cells. It exists during embryogenesis in the blood island region of the yolk sac<sup>20</sup>, which is therefore the earliest site of hematopoiesis and vasculogenesis. Until recently, vasculogenesis has been thought to be restricted to the yolk sac and the early embryogenesis. However, novel observations have revealed in adulthood a situation consistent with vasculogenesis: endothelial cells derived from angioblasts or "hemangioblasts" previously isolated from peripheral blood or bone marrow are incorporated into sites of neovascularization in physiological and pathological conditions<sup>21-25</sup>. In addition, the number of these endothelial cell progenitors increases in the peripheral blood during tissue ischemia or following the administration of VEGF or GM-CSF, a cytokine known to induce mobilization of hematopoietic stem cells from the bone marrow into the peripheral blood<sup>24,25</sup>. Recent studies in humans, dogs, rats, rabbits and mice have indeed indicated the presence of endothelial precursor cells (EPCs) in bone marrow and peripheral blood during adult life which can be mobilized and incorporated into newly formed vessels or are involved in endothelialization of implants<sup>23,26-32</sup>. Interestingly, in all these experiments, endothelial cell progenitors are isolated together with other hematopoietic stem cells by using antibodies directed against hematopoietic stem cell antigens. PROM-1 is expressed on lineage non-committed stem and progenitor cells but not on mature peripheral blood cells and umbilical vein derived endothelial cells<sup>2</sup>. CD34<sup>+</sup> cells co-

expressing VEGFR-2 and PROM-1, have been isolated from peripheral blood, cord blood, fetal liver and bone marrow. When grown in the presence of VEGF and FGF-2 or the cytokine stem cell growth factor (SCGF), these cells give rise to endothelial cells, thus suggesting that this subset of CD34<sup>+</sup>, VEGFR-2<sup>+</sup> & CD133<sup>+</sup> cells may play a role in neovasculogenesis<sup>3,5,11</sup>. The present invention uses a transgenic mouse deficient in PROM-1 to study the involvement of PROM-1 in several pathological models of angiogenesis. For the sake of clarity the nucleotide sequence of human prominin-1 is designated here as SEQ ID NO: 1 and the amino acid sequence of human prominin-1 is designated as SEQ ID NO: 2. The present invention shows that inhibitors of prominin-1 can be used in therapeutic applications for the prevention of pathological angiogenesis.

Thus the invention provides in one embodiment the use of a molecule which comprises a region specifically binding to prominin-1 (SEQ ID NO: 2) or nucleic acids encoding prominin-1 (SEQ ID NO: 1), for the manufacture of a medicament to treat pathological angiogenesis.

According to the invention molecules that comprise a region specifically binding to prominin-1 or nucleic acids encoding prominin-1 which can be used for the manufacture of a medicament to treat pathological angiogenesis can be chosen from the list comprising an antibody or any fragment thereof binding to prominin-1, a (synthetic) peptide, a protein, a small molecule specifically binding to prominin-1 or nucleic acids encoding prominin-1 or a regulatory region (e.g. a promoter region) of prominin-1, RNA aptamers against prominin-1, a ribozyme against nucleic acids encoding prominin-1, anti-sense nucleic acids hybridising with nucleic acids encoding prominin-1 and small interference RNA's (siRNA) against prominin-1.

The wording 'a molecule which comprises a region specifically binding to prominin-1 or nucleic acids encoding prominin-1' relates (1) on the one hand to molecules binding to nucleic acids encoding prominin-1 or to regulatory genetic regions of prominin-1, said molecules inhibit the gene expression of prominin-1 (thus the inhibition of prominin-1 transcription and/or translation of a gene transcript (mRNA) of prominin-1 and (2) on the other hand to molecules that inhibit the activity of the prominin-1 protein. The inhibition of gene expression can be measured conveniently by methods known in the art such as for example RT-PCR analysis of the prominin-1 transcript or for example western blot analysis of the prominin-1 protein, said inhibition is preferably at least 20%, 30%, 40%, 50%, 60%, 70%, 80%, 90% or even higher. Measurement of molecules that bind to the

prominin-1 protein and inhibit the activity of prominin-1 can for example be carried out by various methods for determining pathological angiogenesis as described in the examples of the present invention. Said inhibition of prominin-1 activity is preferably at least 20%, 30%, 40%, 50%, 60%, 70%, 80%, 90% or even higher. Thus in another

5 embodiment the invention provides the use of a molecule that inhibits the expression and/or activity of prominin-1 for the manufacture of a medicament for treatment of pathological angiogenesis. In the latter embodiment activity relates to the gene product (the protein) and expression relates to the gene: mRNA formation and/or translation of the mRNA of prominin-1.

10 The term 'antibody' or 'antibodies' relates to an antibody characterized as being specifically directed against prominin-1 or any functional derivative thereof, with said antibodies being preferably monoclonal antibodies; or an antigen-binding fragment thereof, of the F(ab')<sub>2</sub>, F(ab) or single chain Fv type, or any type of recombinant antibody derived thereof. These antibodies of the invention, including specific polyclonal antisera

15 prepared against prominin-1 or any functional derivative thereof, have no cross-reactivity to others proteins. The monoclonal antibodies of the invention can for instance be produced by any hybridoma liable to be formed according to classical methods from splenic cells of an animal, particularly of a mouse or rat immunized against prominin-1 or any functional derivative thereof, and of cells of a myeloma cell line, and to be selected

20 by the ability of the hybridoma to produce the monoclonal antibodies recognizing prominin-1 or any functional derivative thereof which have been initially used for the immunization of the animals. The monoclonal antibodies according to this embodiment of the invention may be humanized versions of the mouse monoclonal antibodies made by means of recombinant DNA technology, departing from the mouse and/or human

25 genomic DNA sequences coding for H and L chains or from cDNA clones coding for H and L chains. Alternatively the monoclonal antibodies according to this embodiment of the invention may be human monoclonal antibodies. Such human monoclonal antibodies are prepared, for instance, by means of human peripheral blood lymphocytes (PBL) repopulation of severe combined immune deficiency (SCID) mice as described in

30 PCT/EP 99/03605 or by using transgenic non-human animals capable of producing human antibodies as described in US patent 5,545,806. Also fragments derived from these monoclonal antibodies such as Fab, F(ab')<sub>2</sub> and ssFv ("single chain variable fragment"), providing they have retained the original binding properties, form part of the present invention. Such fragments are commonly generated by, for instance, enzymatic

35 digestion of the antibodies with papain, pepsin, or other proteases. It is well known to

the person skilled in the art that monoclonal antibodies, or fragments thereof, can be modified for various uses. The antibodies involved in the invention can be labeled by an appropriate label of the enzymatic, fluorescent, or radioactive type.

In a specific embodiment the antibodies against prominin-1 can be derived from animals of the camelid family. In said family immunoglobulins devoid of light polypeptide chains are found. Heavy chain variable domain sequences derived from camelids are designated as VHH's. "Camelids" comprise old world camelids (*Camelus bactrianus* and *Camelus dromaderius*) and new world camelids (for example *Lama paccos*, *Lama glama* and *Lama vicugna*). EP0656946 describes the isolation and uses of camelid immunoglobulins and is incorporated herein by reference.

Small molecules, e.g. small organic molecules, and other drug candidates can be obtained, for example, from combinatorial and natural product libraries.

Also within the scope of the invention are oligoribonucleotide sequences, that include anti-sense RNA and DNA molecules and ribozymes that function to inhibit the translation of prominin-1 mRNA. Anti-sense RNA and DNA molecules act to directly block the translation of mRNA by binding to targeted mRNA and preventing protein translation. In regard to antisense DNA, oligodeoxyribonucleotides derived from the translation initiation site, e.g., between -10 and +10 regions of the prominin-1 nucleotide sequence, are preferred. Ribozymes are enzymatic RNA molecules capable of catalyzing the specific cleavage of RNA. The mechanism of ribozyme action involves sequence specific hybridization of the ribozyme molecule to complementary target RNA, followed by an endonucleolytic cleavage. Within the scope of the invention are engineered hammerhead motif ribozyme molecules that specifically and efficiently catalyze endonucleolytic cleavage of prominin-1 RNA sequences.

Specific ribozyme cleavage sites within any potential RNA target are initially identified by scanning the target molecule for ribozyme cleavage sites which include the following sequences, GUA, GUU and GUC. Once identified, short RNA sequences of between 15 and 20 ribonucleotides corresponding to the region of the target gene containing the cleavage site may be evaluated for predicted structural features such as secondary structure that may render the oligonucleotide sequence unsuitable. The suitability of candidate targets may also be evaluated by testing their accessibility to hybridization with complementary oligonucleotides, using ribonuclease protection assays.

Both anti-sense RNA and DNA molecules and ribozymes of the invention may be prepared by any method known in the art for the synthesis of RNA molecules. These include techniques for chemically synthesizing oligodeoxyribonucleotides well known in

the art such as for example solid phase phosphoramidite chemical synthesis. Alternatively, RNA molecules may be generated by *in vitro* and *in vivo* transcription of DNA sequences encoding the antisense RNA molecule. Such DNA sequences may be incorporated into a wide variety of vectors which incorporate suitable RNA polymerase promoters such as the T7 or SP6 polymerase promoters. Alternatively, antisense cDNA constructs that synthesize anti-sense RNA constitutively or inducibly, depending on the promoter used, can be introduced stably into cell lines.

In a particular embodiment short interference RNA molecules (siRNA) can be used for the manufacture of a medicament for treatment of pathological angiogenesis. Said interference RNA molecules can be generated based on the genetic sequence of prominin-1 (SEQ ID NO: 1). RNA interference (RNAi) is based on the degradation of particular target sequences by the design of short interference RNA oligo's (siRNA) which recognize the target sequence (here SEQ ID NO: 1) and subsequently trigger their degradation by a poorly understood pathway. In general siRNA duplexes are shorter than 30 nucleotides, because longer stretches of dsRNA activate the PKR pathway in mammalian cells which results in a global a-specific shut-down of protein synthesis. The preparation and gene therapy vectors for the intracellular expression of siRNAs duplexes is disclosed in WO0244321 which is herein incorporated by reference.

In another particular embodiment RNA aptamers can be used for the manufacture of a medicament for treatment of pathological angiogenesis. Said RNA aptamers can be generated against prominin-1 (SEQ ID NO: 2). Recently, RNA aptamers have been used as therapeutic reagents for their ability to disrupt protein function. Selection of aptamers *in vitro* allows rapid isolation of extremely rare RNAs that have high specificity and affinity for specific proteins. Exemplary RNA aptamers are described in U.S. Pat. No. 5,270,163 to Gold et al., Ellington and Szostak, "In vitro Selection of RNA Molecules That Bind Specific Ligands," *Nature* 346:818-822 (1990), and Tuerk and Gold, "Systematic Evolution of Ligands by Exponential Enrichment: RNA Ligands to Bacteriophage T4 DNA Polymerase," *Science* 249:505-510 (1990). Unlike antisense compounds, whose targets are one dimensional lattices, RNA aptamers can bind to the three dimensional surfaces of a protein. Moreover, RNA aptamers can frequently discriminate finely among discrete functional sites of a protein (Gold et al., "Diversity of Oligonucleotide Functions," *Annu. Rev. Biochem.* 64:763-797 (1995)). As research and therapeutic reagents, aptamers not only have the combined advantages of antibodies and small molecular mass drugs, but *in vivo* production of RNA aptamers also can be

controlled genetically Such RNA expressing genes are usually smaller than protein-coding genes and can be inserted into gene therapy vectors.

The term 'pathological angiogenesis' refers to the excessive formation and growth of blood vessels during the maintenance and the progression of several disease states.

- 5 Examples where pathological angiogenesis can occur are blood vessels (atherosclerosis, hemangioma, hemangioendothelioma), bone and joints (rheumatoid arthritis, synovitis, bone and cartilage destruction, osteomyelitis, pannus growth, osteophyte formation, neoplasms and metastasis), skin (warts, pyogenic granulomas, hair growth, Kaposi's sarcoma, scar keloids, allergic oedema, neoplasms), liver, kidney, lung, ear and other epithelia (inflammatory and infectious processes (including hepatitis, glomerulonephritis, pneumonia), asthma, nasal polyps, otitis, transplantation, liver regeneration, neoplasms and metastasis), uterus, ovary and placenta (dysfunctional uterine bleeding (due to intrauterine contraceptive devices), follicular cyst formation, ovarian hyperstimulation syndrome, endometriosis, neoplasms), brain, nerves and eye (retinopathy of prematurity, diabetic retinopathy, choroidal and other intraocular disorders, leukomalacia, neoplasms and metastasis), heart and skeletal muscle due to work overload, adipose tissue (obesity), endocrine organs (thyroiditis, thyroid enlargement, pancreas transplantation), hematopoiesis (AIDS (Kaposi), hematologic malignancies (leukemias, etc.), tumour induced new blood vessels.
- 15
- 20 The term 'medicament to treat' relates to a composition comprising molecules as described above and a pharmaceutically acceptable carrier or excipient (both terms can be used interchangeably) to treat diseases as indicated above. Suitable carriers or excipients known to the skilled man are saline, Ringer's solution, dextrose solution, Hank's solution, fixed oils, ethyl oleate, 5% dextrose in saline, substances that enhance isotonicity and chemical stability, buffers and preservatives. Other suitable carriers include any carrier that does not itself induce the production of antibodies harmful to the individual receiving the composition such as proteins, polysaccharides, polylactic acids, polyglycolic acids, polymeric amino acids and amino acid copolymers. The 'medicament' may be administered by any suitable method within the knowledge of the skilled man. The preferred route of administration is parenterally. In parental administration, the medicament of this invention will be formulated in a unit dosage injectable form such as a solution, suspension or emulsion, in association with the pharmaceutically acceptable excipients as defined above. However, the dosage and mode of administration will depend on the individual. Generally, the medicament is administered so that the protein, peptide, antibody, small molecule, ribozyme, RNA
- 25
- 30
- 35

aptamer, anti-sense nucleic acid or siRNA of the present invention is given at a dose between 1  $\mu\text{g/kg}$  and 10 mg/kg, more preferably between 10  $\mu\text{g/kg}$  and 5 mg/kg, most preferably between 0.1 and 2 mg/kg. Preferably, it is given as a bolus dose. Continuous infusion may also be used and includes continuous subcutaneous delivery  
5 via an osmotic minipump. If so, the medicament may be infused at a dose between 5 and 20  $\mu\text{g/kg/minute}$ , more preferably between 7 and 15  $\mu\text{g/kg/minute}$ .

In another embodiment antibodies or functional fragments thereof can be used for the manufacture of a medicament for the treatment of the above mentioned disorders. As a non-limiting example there are the antibodies described in US 5,843,633. In a specific  
10 embodiment said antibodies are humanized (Rader et al., 2000, J. Biol. Chem. 275, 13668) and more specifically human antibodies are used to manufacture a medicament to treat pathological angiogenesis. In yet another specific embodiment antibodies derived from camelids are used to manufacture a medicament to treat pathological angiogenesis.

Another aspect of administration for treatment is the use of gene therapy to deliver the above mentioned anti-sense gene or functional parts of the prominin-1 gene or a ribozyme directed against the prominin-1 mRNA or a functional part thereof or RNA aptamers or siRNAs. Gene therapy means the treatment by the delivery of therapeutic nucleic acids to patient's cells. This is extensively reviewed in Lever and Goodfellow  
15 1995; Br. Med Bull., 51, 1-242; Culver 1995; Ledley, F.D. 1995. Hum. Gene Ther. 6, 1129. To achieve gene therapy there must be a method of delivering genes to the patient's cells and additional methods to ensure the effective production of any therapeutic genes. There are two general approaches to achieve gene delivery; these are non-viral delivery and virus-mediated gene delivery.

25 The invention also provides methods for identifying compounds or molecules which bind on prominin-1 and prevent or suppress pathological angiogenesis. With "suppression" it is understood that said suppression can occur for at least 20%, 30%, 30%, 50%, 60%, 70%, 80%, 90% or even 100%.

30 Thus in another embodiment the invention provides a method to identify molecules that comprise a region that specifically binds to prominin-1 comprising: (1) exposing prominin-1 or nucleic acids encoding prominin-1 to at least one molecule whose ability to suppress or prevent pathological angiogenesis is sought to be determined, (2) determining binding or hybridising of said molecule(s) to prominin-1 or nucleic acids



encoding prominin-1, and (3) monitoring said pathological angiogenesis when administering said molecules as a medicament.

The latter method is also referred to as 'drug screening assay' or 'bioassay' and typically include the step of screening a candidate/test compound or agent for the ability to interact with prominin-1. Candidate compounds or agents, which have this ability, can be used as drugs to combat or prevent pathological conditions of angiogenesis. Candidate/test compounds are described herein before and are for example RNA aptamers, others are small molecules, e.g. small organic molecules, and other drug candidates can be obtained, for example, from combinatorial and natural product libraries as described above. Typically, the assays are cell-free assays which include the steps of combining prominin-1 and a candidate/test compound (molecule), e.g., under conditions which allow for interaction of (e.g. binding of) the candidate/test compound with prominin-1 to form a complex, and detecting the formation of a complex, in which the ability of the candidate compound to interact with prominin-1 is indicated by the presence of the candidate compound in the complex. Formation of complexes between prominin-1 and the candidate compound can be quantitated, for example, using standard immunoassays. The prominin-1 employed in such a test may be free in solution, affixed to a solid support, borne on a cell surface, or located extracellularly or even intracellularly.

To perform the above described drug screening assays, it is feasible to immobilize prominin-1 or its (their) target molecule(s) to facilitate separation of complexes from uncomplexed forms of one or both of the proteins, as well as to accommodate automation of the assay. Interaction (e.g., binding of) of prominin-1 to a target molecule, can be accomplished in any vessel suitable for containing the reactants.

Examples of such vessels include microtitre plates, test tubes, and microcentrifuge tubes. In one embodiment, a fusion protein can be provided which adds a domain that allows the protein to be bound to a matrix. For example, prominin-1-His tagged can be adsorbed onto Ni-NTA microtitre plates, or prominin-1-ProtA fusions adsorbed to IgG, which are then combined with the cell lysates (e.g., <sup>35</sup>S-labeled) and the candidate compound, and the mixture incubated under conditions conducive to complex formation (e.g., at physiological conditions for salt and pH). Following incubation, the plates are washed to remove any unbound label, and the matrix immobilized and radiolabel determined directly, or in the supernatant after the complexes are dissociated. Alternatively, the complexes can be dissociated from the matrix, separated by SDS-PAGE, and the level of prominin-1-binding protein found in the bead fraction quantitated

from the gel using standard electrophoretic techniques. Other techniques for immobilizing protein on matrices can also be used in the drug screening assays of the invention. For example, prominin-1 can be immobilized utilizing conjugation of biotin and streptavidin. Biotinylated prominin-1 can be prepared from biotin-NHS (N-hydroxy-succinimide) using techniques well known in the art (e.g., biotinylation kit, Pierce Chemicals, Rockford, Ill.), and immobilized in the wells of streptavidin-coated 96 well plates (Pierce Chemical). Another technique for drug screening which provides for high throughput screening of compounds having suitable binding affinity to prominin-1 is described in detail in "Determination of Amino Acid Sequence Antigenicity" by Geysen HN, WO 84/03564, published on 13/09/84. In summary, large numbers of different small peptide test compounds are synthesized on a solid substrate, such as plastic pins or some other surface. The protein test compounds are reacted with fragments of prominin-1 and washed. Bound prominin-1 is then detected by methods well known in the art. Purified prominin-1 can also be coated directly onto plates for use in the aforementioned drug screening techniques. Alternatively, non-neutralizing antibodies can be used to capture the peptide and immobilize it on a solid support. This invention also contemplates the use of competitive drug screening assays in which neutralizing antibodies capable of binding prominin-1 specifically compete with a test compound for binding prominin-1. In this manner, the antibodies can be used to detect the presence of any protein, which shares one or more antigenic determinants with prominin-1.

## Examples

### 1. Generation of a prominin-1 knock-out mice

To study the *in vivo* role of PROM-1, PROM-1 (prominin-1) deficient mice were generated. Targeted inactivation of the PROM-1 gene was achieved by deletion of exon 2 (containing the start codon). Briefly, a genomic BAC (bacterial artificial chromosome) containing the murine PROM-1<sup>1</sup> was obtained from Research Genetics Inc (Huntsville, AL) after screening by PCR and hybridization. Mapping of the murine PROM-1 homologue gene revealed that the first exon, which is 79 bp long, is separated from the second exon by an approximately 8 kb intron. It is the second exon (376 bp long) that contains the startcodon ATG. A BamHI fragment of 11.5 kb containing exon 2 was subcloned into the pUC18 plasmid. A targeting vector for inactivation of the PROM-1 gene, pPNT.PROM-1<sup>null</sup>, was constructed consisting of, from 5' to 3': 1.2 kb of 5' homology comprising the end of intron 1; a *loxP*-flanked *neomycin* gene; 5.5 kb from

intron 2 as 3'-homology; and a *thymidine kinase* selection cassette outside of the regions of homology for counterselection against random integration events. The integrity of the construct was verified by restriction digestion and sequencing. The linearized targeting vector *pPNT.PROM-1<sup>null</sup>* was electroporated in R1 ES cells and targeted clones were identified by appropriate Southern blot analysis and used for morula aggregation to generate PROM-1 deficient chimeric and germline mice. PROM-1 deficient mice were born at the expected Mendelian frequency (~25% of 450 offspring from PROM-1 heterozygous breeding pairs). They appeared healthy and were fertile, irrespective of their genetic background (backgrounds tested: 50% Swiss/50% 129, 100% 129, 50% C57Bl6/50% 129). We anticipated that PROM-1 might play a crucial role in hematopoiesis implying that the PROM-1<sup>-/-</sup> embryo would die *in utero* either after the appearance of the primitive hematopoiesis (7.5 days post coitum, site: yolk sac) or at the emergence of the definitive hematopoiesis (12.5 days post coitum, site: fetal liver). Surprisingly, however, embryonic development in PROM-1<sup>-/-</sup> mice was normal. PROM-1<sup>-/-</sup> embryos were not rescued by maternal PROM-1, as PROM-1<sup>-/-</sup> embryos, sired by PROM-1<sup>+/-</sup> as well as by PROM-1<sup>-/-</sup> breeding pairs, developed normally. Also postnatal physiological vascular development seemed normal since no vascular defects could be observed in the heart (capillary density is  $5810 \pm 154$  in PROM-1<sup>+/+</sup> pups versus  $5394 \pm 179$  in PROM-1<sup>-/-</sup> pups,  $n=3$ ;  $p=NS$ ), kidneys, lungs and skeletal muscle during postnatal growth in PROM-1<sup>-/-</sup> mice.

## 2. Impaired pathological angiogenesis and/or vasculogenesis in prominin-1 knock-out mice

In order to study the role of PROM-1 in pathological conditions of angiogenesis PROM-1<sup>-/-</sup> mice and their wild-type littermates are subjected to various murine models of pathological blood vessel formation.

### 2.1 Ischemic retinopathy

PROM-1<sup>-/-</sup> mice and their wild-type littermates were subjected to a mouse model of ischemic retinopathy. In this hyperoxia-induced retinopathy model, neonatal mice (with an immature retinal vasculature) are exposed to hyperoxia, resulting in obliteration of the developing blood vessels supplying oxygen to the retina. When the mice are then returned to normoxia, the retina, distal to the occluded vessels, becomes ischemic,

inducing VEGF production and ultimately resulting in reproducible and quantifiable proliferative retinal neovascularization (33, 34). This model, which mimicks to a certain extent the vascular response during retinopathy of prematurity or diabetic retinopathy, may be useful to test the efficacy of (anti)-angiogenic molecules (Pierce EA et al (1995) Proc. Natl. Acad. Sci. 92(3)905-9). Mouse pups of 7 days (P7) together with their mothers, are subjected to hyperoxia (75% oxygen) in specially designed oxygen chambers for 5 days, without opening the cages. On P12, the animals are returned to room air until P17, when the retinas are assessed for maximal neovascular response. Anaesthetized mice are perfused through the left ventricle with 1 ml of phosphate buffered saline containing 50 mg of  $2 \times 10^6$  molecular weight fluorescein-dextran. The eyes are removed and fixed in 4% paraformaldehyde for 3 (right eye) or 24 (left eye) hrs. Of the right eyes, lenses are removed and peripheral retinas cut to allow flat mounting with glycerol-gelatin. The flat mounted retinas are analyzed by fluorescence microscopy. The left eyes are embedded in paraffin and serial 6  $\mu\text{m}$  sections are cut sagittally throughout the cornea, parallel to the optic nerve, and stained with hematoxylin-eosin. The proliferative neovascular response is quantified by counting the number of new vessels (= tufts) and the number of endothelial cells extending from the internal limiting membrane of the retina into the vitreum on the stained sagittal cross-sections. The angiographic technique using fluorescein-dextran perfusion is used in conjunction with this counting method for rapid screening of retinas or as an alternative grading system for quantitative evaluation. Loss of prominin-1 significantly protected mice against intra-vitreous neovascularization, as evaluated by counting the number of neovascular tufts and endothelial cells (EC) in the vitreous cavity ( $n=15$ ;  $p<0.001$ )

	N° of tufts in vitreous cavity (per 10 retinal sections)	N° of EC in vitreous cavity (per 10 retinal sections)
PROM-1 <sup>-/-</sup> <sup>+/+</sup> pups (n = 15)	157.1 $\pm$ 13.6	286.0 $\pm$ 45.1
PROM-1 <sup>-/-</sup> pups (n = 15)	72.5 $\pm$ 14.6	106.2 $\pm$ 22.6

## 2.2 Corneal micropocket assay

Hydron pellets containing an angiogenic substance (like bFGF or VEGF) are implanted into the corneal stroma adjacent to the temporal limbus. This induces neovascularization of the avascular corneal stroma from day 3 to day 8 after implantation, without substantial corneal edema or inflammation. Like the retinal hypoxia model, it gives a predictable, persistent and aggressive neovascular response, which is dependent on direct stimulation of blood vessels rather than on indirect stimulation by the induction of inflammation<sup>35</sup>. The mouse corneal micropocket assay was performed as previously described<sup>36</sup>. Hydron-coated sucralfate pellets containing 300 ng of VEGF<sub>165</sub> were positioned 1 mm from the corneal limbus. Mice deficient for PROM-1 showed a reduced angiogenic response. The length of the newly formed vessels ( $0.93 \pm 0.12$  mm in PROM-1<sup>+/+</sup> mice versus  $0.70 \pm 0.03$  mm in PROM-1<sup>-/-</sup> mice, n=6; p<0.005) as well as the circumferential neovascularity ( $6.23 \pm 0.55$  mm in PROM-1<sup>+/+</sup> mice versus  $3.60 \pm 0.32$  mm in PROM-1<sup>-/-</sup> mice, n=6; p<0.005) and the integrated optical density of the vessel area ( $497 \pm 100$  in PROM-1<sup>+/+</sup> mice versus  $196 \pm 27$  in PROM-1<sup>-/-</sup> mice, n=6; p<0.05) were significantly lower in the PROM-1 mice. Moreover, WT bone marrow transplantation into PROM-1 deficient mice rescued the impaired angiogenic response. The length of the newly formed vessels ( $0.57 \pm 0.03$  mm in PROM-1<sup>+/+</sup> mice versus  $0.57 \pm 0.03$  mm in PROM-1<sup>-/-</sup> mice, n=6; p<0.005) as well as the circumferential neovascularity ( $3.52 \pm 0.26$  mm in PROM-1<sup>+/+</sup> mice versus  $3.29 \pm 0.22$  mm in PROM-1<sup>-/-</sup> mice, n=6; p=NS) were identical in both PROM-1 deficient and WT mice after WT bone marrow transplantation.

## 2.3 Model of skin wound healing

Vascular remodeling was studied in a model of skin wound healing as described before<sup>37,38</sup>. For skin wounding, a standardised 15 mm full-thickness skin incision was made on the back of the mice, taking care not to damage the underlying muscle. Wound healing was quantified by daily measuring the width and the length of the wound. New blood vessel formation was analysed on skin sections harvested four days after wounding. Wound healing was significantly impaired in the PROM-1 deficient mice. Both genotypes contained comparable densities of vessels in unwounded skin. However, the number of capillaries infiltrating the wound ( $185.8 \pm 11.1$  vessels/mm<sup>2</sup> in PROM-1<sup>+/+</sup> mice versus  $135.0 \pm 12.7$  in PROM-1<sup>-/-</sup> mice, n=5; p<0.05), as well as the number of

smooth muscle-coated vessels in the wounded area ( $58.2 \pm 10$  vessels/mm<sup>2</sup> in PROM-1<sup>+/+</sup> mice versus  $28.6 \pm 4.022$  in PROM-1<sup>-/-</sup> mice, n=5; p<0.05) were significantly reduced in PROM-1 deficient mice.

#### 5 2.4 Matrigel assay

In-growth of capillaries was studied in a matrigel assay performed as described<sup>39</sup>. The angiogenic response in the matrigel of PROM-1<sup>-/-</sup> mice seemed somewhat lower as measured by the hemoglobin content per matrigel implant ( $137.0 \pm 20.4$  µg/ml in PROM-1<sup>+/+</sup> mice *versus*  $112.1 \pm 17.6$  µg/ml in PROM-1<sup>-/-</sup> mice; n=15; p= NS).

10 Histological sections of matrigel were then analysed for infiltration of leukocytes and for blood vessel formation after staining for inflammatory cells (CD45) and endothelial cells (CD31), respectively. The number of infiltrating leukocytes did not seem to differ but a reduced blood vessel formation (CD31 positive endothelial cells) was noticed in the matrigel implanted in PROM-1 deficient animals (% of CD31 positive area in matrigel:  
15  $0.55 \pm 0.08\%$  in PROM-1<sup>+/+</sup> mice *versus*  $0.26 \pm 0.06\%$  in PROM-1<sup>-/-</sup> mice; n=5; p<0.05).

#### 2.5 Myocardial infarction model

Myocardial infarction was performed by ligation of the LAD as described<sup>40</sup>. After 4 to 7 days, infarcted hearts were used for histological analysis or for immunostaining of  
20 thrombomodulin (endothelial cells) or smooth muscle alpha-actin (smooth muscle cells)  
<sup>38</sup>. Morphometric analysis and counting of immunoreactive cells was performed using an image analysis system with KS300 software (Zeiss, Brussels, Belgium). No differences were observed in the number of capillaries at 4 ( $490.9 \pm 65.4$  vessels/mm<sup>2</sup> in PROM-1<sup>+/+</sup> mice *versus*  $493.3 \pm 87.7$  vessels/mm<sup>2</sup> in PROM-1<sup>-/-</sup> mice; n=3; p= NS) or  
25 7 days ( $510.6 \pm 28.3$  vessels/mm<sup>2</sup> in PROM-1<sup>+/+</sup> mice *versus*  $507.8 \pm 24.6$  vessels/mm<sup>2</sup> in PROM-1<sup>-/-</sup> mice; n=10; p= NS) after ligation or in the number of SMC covered vessels at 4 ( $20.3 \pm 3.2$  vessels/mm<sup>2</sup> in PROM-1<sup>+/+</sup> mice *versus*  $23.3 \pm 6.73$  vessels/mm<sup>2</sup> in PROM-1<sup>-/-</sup> mice; n=3; p= NS) or at 7 days ( $87.6 \pm 14.3$  vessels/mm<sup>2</sup> in PROM-1<sup>+/+</sup> mice *versus*  $76.4 \pm 12.2$  vessels/mm<sup>2</sup> in PROM-1<sup>-/-</sup> mice; n=10; p= NS) in  
30 the infarcted area of hearts of PROM-1 deficient mice and wild-type littermates. However, a clear significant difference was observed in the number of infiltrating macrophages at 7 days after ligation (% of Mac3 positive area:  $3.75 \pm 0.77$  % in PROM-1<sup>+/+</sup> mice *versus*  $1.62 \pm 0.42$  % in PROM-1<sup>-/-</sup> mice; n=10; p< 0.05).

### 2.6 Hind limb ischemia model

Hind limb ischemia is induced as described <sup>41</sup>. Unilateral right or bilateral ligations of the femoral artery and vein (proximal to the popliteal artery) and the cutaneous vessels branching from the caudal femoral artery side branch is performed and two superficial preexisting collateral arterioles, connecting the femoral and saphenous artery, will be used for analysis. Genetic consequences on post-ischemic revascularization is determined 14 days after ligation, using vascular morphological (histological evaluation of capillary density and SMC-coated vessel density, histological evaluation of myocyte necrosis and regeneration), perfusional (fluorescent microspheres, laser Doppler imaging), and functional (graded treadmill exercise or swim endurance exercise) analyses.

### 2.7 Tumor models

The role of PROM-1 is also tested in tumor models. The following mouse models are operational and are used to analyze tumor angiogenesis in vivo: 1) subcutaneous injection of ras-transformed fibroblasts in athymic nude (nu/nu) mice, 2) subcutaneous injection of Lewis lung carcinoma cells in syngenic C57Bl6 hosts, and 3) subcutaneous inoculation of rat C6 glioma cells of athymic nude (nu/nu) mice. Five to twenty million of tumor cells are inoculated in the mice and tumor growth is followed up to 30 days. Tumors are measured with calipers and tumor volumes calculated using the formula  $[\pi/6 \times (w1 \times w2 \times w2)]$ , where "w1" and "w2" represent the largest and smallest tumor diameter, respectively. Tumor vessel density and size are determined on tissue sections using immunohistochemistry for visualization of endothelial cells (CD-31), in combination with quantitative morphometry of vascular densities and patterning. If necessary, intratumor flow is determined using colored microspheres to quantitate flow across the entire tumor. When WT RAS transformed fibroblasts were injected in PROM-1 deficient and WT nude mice, no difference in tumor weight was seen (tumor weight after 14 days:  $0.9 \pm 0.1$  g in PROM-1<sup>+/+</sup> hosts versus  $1.1 \pm 0.3$  g in PROM-1<sup>-/-</sup> hosts; n=7; p< 0.05). Blood vessel analysis is being performed. Remarkably, the number of infiltrating leukocytes was significantly reduced in the tumors grown in the PROM-1 deficient mice.

### 2.8 LPS induced venous thrombosis in footpad

To study whether PROM-1 is important in inflammatory processes, a chronic

inflammation footpad assay was used. Endotoxin (20 $\mu$ l, E. coli lipopolysaccharide, 5 and 50  $\mu$ g/ml) was injected into the right footpad of both PROM-1 deficient and WT mice as described (Carmeliet, P. et al (1993) J. Clin. Invest 6: 2756-60). Saline is injected into the left footpad as a control. After 5 days, mice were sacrificed and both right and left  
5 footpad were measured with callipers, excised and fixed in 1% paraformaldehyde for 24 hours. Subsequently, footpads were embedded in paraffin and sectioned. Veins are scored on haematoxylin and eosin stained sections for the presence of thrombi. Five days after injecting 50  $\mu$ g/ml of endotoxin, a decrease in footpad thickness was observed in the PROM-1 deficient compared to their WT controls.

10

These data clearly indicate a role of AC 133 in pathological vasculogenesis and/or angiogenesis and implicate the use of inhibitors of PROM-1 in therapeutic strategies to inhibit blood vessel formation in various pathological disorders.



References

1. Miraglia S, Godfrey W, Yin AH, Atkins K, Warnke R, Holden JT, Bray RA, Waller EK, Buck DW. A novel five-transmembrane hematopoietic stem cell antigen: isolation, characterization, and molecular cloning. Blood 1997;90(12):5013-21.
2. Yin AH, Miraglia S, Zanjani ED, Almeida-Porada G, Ogawa M, Leary AG, Olweus J, Kearney J, Buck DW. PROM-1, a novel marker for human hematopoietic stem and progenitor cells. Blood 1997;90(12):5002-12.
3. Gehling UM, Ergun S, Schumacher U, Wagener C, Pantel K, Otte M, Schuch G, Schafhausen P, Mende T, Kilic N and others. In vitro differentiation of endothelial cells from PROM-1-positive progenitor cells. Blood 2000;95(10):3106-12.
4. Boyer M, Townsend LE, Vogel LM, Falk J, Reitz-Vick D, Trevor KT, Villalba M, Bendick PJ, Glover JL. Isolation of endothelial cells and their progenitor cells from human peripheral blood. J Vasc Surg 2000;31(1 Pt 1):181-9.
5. Quirici N, Soligo D, Caneva L, Servida F, Bossolasco P, Deliliers GL. Differentiation and expansion of endothelial cells from human bone marrow CD133(+) cells. Br J Haematol 2001;115(1):186-94.
6. Buhring HJ, Seiffert M, Bock TA, Scheduling S, Thiel A, Scheffold A, Kanz L, Brugger W. Expression of novel surface antigens on early hematopoietic cells. Ann N Y Acad Sci 1999;872:25-38; discussion 38-9.
7. Horn PA, Tesch H, Staib P, Kube D, Diehl V, Voliotis D. Expression of PROM-1, a novel hematopoietic precursor antigen, on acute myeloid leukemia cells. Blood 1999;93(4):1435-7.
8. Kratz-Albers K, Zuhlsdorf M, Leo R, Berdel WL, Buchner T, Serve H. Expression of a PROM-1, a novel stem cell marker, on human leukemic blasts lacking CD34-antigen and on a human CD34+ leukemic line:MUTZ-2. Blood 1998;92(11):4485-7.
9. Buhring HJ, Seiffert M, Marxer A, Weiss B, Faul C, Kanz L, Brugger W. PROM-1 antigen expression is not restricted to acute myeloid leukemia blasts but is also found on acute lymphoid leukemia blasts and on a subset of CD34+ B-cell precursors. Blood 1999;94(2):832-3.
10. Corbeil D, Roper K, Hannah MJ, Hellwig A, Huttner WB. Selective localization of the polytopic membrane protein prominin in microvilli of epithelial cells - a combination of apical sorting and retention in plasma membrane protrusions. J Cell Sci 1999;112(Pt 7):1023-33.

11. Peichev M, Naiyer AJ, Pereira D, Zhu Z, Lane WJ, Williams M, Oz MC, Hicklin DJ, Witte L, Moore MA and others. Expression of VEGFR-2 and PROM-1 by circulating human CD34(+) cells identifies a population of functional endothelial precursors. *Blood* 2000;95(3):952-8.
- 5 12. Rafii S. Circulating endothelial precursors: mystery, reality, and promise. *J Clin Invest* 2000;105(1):17-9.
13. Uchida N, Buck DW, He D, Reitsma MJ, Masek M, Phan TV, Tsukamoto AS, Gage FH, Weissman IL. Direct isolation of human central nervous system stem cells. *Proc Natl Acad Sci U S A* 2000;97(26):14720-5.
- 10 14. Corbeil D, Roper K, Hellwig A, Tavian M, Miraglia S, Watt SM, Simmons PJ, Peault B, Buck DW, Huttner WB. The human PROM-1 hematopoietic stem cell antigen is also expressed in epithelial cells and targeted to plasma membrane protrusions. *J Biol Chem* 2000;275(8):5512-20.
- 15 15. Yu Y, Flint A, Dvorin EL, Bischoff J. PROM-1-2, a Novel Isoform of Human PROM-1 Stem Cell Antigen. *J Biol Chem* 2002;277(23):20711-6.
16. Maw MA, Corbeil D, Koch J, Hellwig A, Wilson-Wheeler JC, Bridges RJ, Kumaramanickavel G, John S, Nancarrow D, Roper K and others. A frameshift mutation in prominin (mouse)-like 1 causes human retinal degeneration. *Hum Mol Genet* 2000;9(1):27-34.
- 20 17. Corbeil D, Roper K, Fargeas CA, Joester A, Huttner WB. Prominin: a story of cholesterol, plasma membrane protrusions and human pathology. *Traffic* 2001;2(2):82-91.
18. Corbeil D, Roper K, Weigmann A, Huttner WB. PROM-1 hematopoietic stem cell antigen: human homologue of mouse kidney prominin or distinct member of a novel protein family? *Blood* 1998;91(7):2625-6.
- 25 19. Miraglia S, Godfrey W, Buck D. A response to PROM-1 hematopoietic stem cell antigen: human homologue of mouse kidney prominin or distinct member of a novel protein family? *Blood* 1998;91(11):4390-1.
20. Risau W. Mechanisms of angiogenesis. *Nature* 1997;386(6626):671-4.
- 30 21. Asahara T, Murohara T, Sullivan A, Silver M, van der Zee R, Li T, Witzenbichler B, Schatteman G, Isner JM. Isolation of putative progenitor endothelial cells for angiogenesis. *Science* 1997;275(5302):964-7.
22. Shi Q, Rafii S, Wu MH, Wijelath ES, Yu C, Ishida A, Fujita Y, Kothari S, Mohle R, Sauvage LR and others. Evidence for circulating bone marrow-derived endothelial cells. *Blood* 1998;92(2):362-7.
- 35

23. Takahashi T, Kalka C, Masuda H, Chen D, Silver M, Kearney M, Magner M, Isner JM, Asahara T. Ischemia- and cytokine-induced mobilization of bone marrow-derived endothelial progenitor cells for neovascularization. *Nat Med* 1999;5(4):434-8.
- 5 24. Asahara T, Masuda H, Takahashi T, Kalka C, Pastore C, Silver M, Kearne M, Magner M, Isner JM. Bone marrow origin of endothelial progenitor cells responsible for postnatal vasculogenesis in physiological and pathological neovascularization. *Circ Res* 1999;85(3):221-8.
- 10 25. Asahara T, Takahashi T, Masuda H, Kalka C, Chen D, Iwaguro H, Inai Y, Silver M, Isner JM. VEGF contributes to postnatal neovascularization by mobilizing bone marrow-derived endothelial progenitor cells. *Embo J* 1999;18(14):3964-72.
- 15 26. Bhattacharya V, McSweeney PA, Shi Q, Bruno B, Ishida A, Nash R, Storb RF, Sauvage LR, Hammond WP, Wu MH. Enhanced endothelialization and microvessel formation in polyester grafts seeded with CD34(+) bone marrow cells. *Blood* 2000;95(2):581-5.
- 20 27. Kalka C, Masuda H, Takahashi T, Kalka-Moll WM, Silver M, Kearney M, Li T, Isner JM, Asahara T. Transplantation of ex vivo expanded endothelial progenitor cells for therapeutic neovascularization. *Proc Natl Acad Sci U S A* 2000;97(7):3422-7.
- 25 28. Schatteman GC, Hanlon HD, Jiao C, Dodds SG, Christy BA. Blood-derived angioblasts accelerate blood-flow restoration in diabetic mice. *J Clin Invest* 2000;106(4):571-8.
- 30 29. Kawamoto A, Gwon HC, Iwaguro H, Yamaguchi JI, Uchida S, Masuda H, Silver M, Ma H, Kearney M, Isner JM and others. Therapeutic potential of ex vivo expanded endothelial progenitor cells for myocardial ischemia. *Circulation* 2001;103(5):634-7.
- 30 30. Kocher AA, Schuster MD, Szabolcs MJ, Takuma S, Burkhoff D, Wang J, Homma S, Edwards NM, Itescu S. Neovascularization of ischemic myocardium by human bone-marrow-derived angioblasts prevents cardiomyocyte apoptosis, reduces remodeling and improves cardiac function. *Nat Med* 2001;7(4):430-6.
31. Vasa M, Fichtlscherer S, Aicher A, Adler K, Urbich C, Martin H, Zeiher AM, Dimmeler S. Number and migratory activity of circulating endothelial progenitor cells inversely correlate with risk factors for coronary artery disease. *Circ Res* 2001;89(1):E1-7.

32. Gill M, Dias S, Hattori K, Rivera ML, Hicklin D, Witte L, Girardi L, Yurt R, Himel H, Rafii S. Vascular trauma induces rapid but transient mobilization of VEGFR2(+)PROM-1(+) endothelial precursor cells. *Circ Res* 2001;88(2):167-74.
33. Smith LE, Wesolowski E, McLellan A, Kostyk SK, D'Amato R, Sullivan R, D'Amore PA. Oxygen-induced retinopathy in the mouse. *Invest Ophthalmol Vis Sci* 1994;35(1):101-11.
34. Wesolowski E, Smith LE. Effect of light on oxygen-induced retinopathy in the mouse. *Invest Ophthalmol Vis Sci* 1994;35(1):112-9.
35. Kenyon BM, Voest EE, Chen CC, Flynn E, Folkman J, D'Amato RJ. A model of angiogenesis in the mouse cornea. *Invest Ophthalmol Vis Sci* 1996;37(8):1625-32.
36. Asahara T, Chen D, Takahashi T, Fujikawa K, Kearney M, Magner M, Yancopoulos GD, Isner JM. Tie2 receptor ligands, angiopoietin-1 and angiopoietin-2, modulate VEGF- induced postnatal neovascularization. *Circ Res* 1998;83(3):233-40.
37. Frank S, Hubner G, Breier G, Longaker MT, Greenhalgh DG, Werner S. Regulation of vascular endothelial growth factor expression in cultured keratinocytes. Implications for normal and impaired wound healing. *J Biol Chem* 1995;270(21):12607-13.
38. Carmeliet P, Moons L, Luttun A, Vincenti V, Compernelle V, De Mol M, Wu Y, Bono F, Devy L, Beck H and others. Synergism between vascular endothelial growth factor and placental growth factor contributes to angiogenesis and plasma extravasation in pathological conditions. *Nat Med* 2001;7(5):575-83.
39. Passaniti A, Taylor RM, Pili R, Guo Y, Long PV, Haney JA, Pauly RR, Grant DS, Martin GR. A simple, quantitative method for assessing angiogenesis and antiangiogenic agents using reconstituted basement membrane, heparin, and fibroblast growth factor. *Lab Invest* 1992;67(4):519-28.
40. Heymans S, Luttun A, Nuyens D, Theilmeier G, Creemers E, Moons L, Dyspersin GD, Cleutjens JP, Shipley M, Angellilo A and others. Inhibition of plasminogen activators or matrix metalloproteinases prevents cardiac rupture but impairs therapeutic angiogenesis and causes cardiac failure. *Nat Med* 1999;5(10):1135-42.
41. Luttun A, Tjwa M, Moons L, Wu Y, Angelillo-Scherrer A, Liao F, Nagy JA, Hooper A, Priller J, De Klerck B and others. Revascularization of ischemic tissues by

- PIGF treatment, and inhibition of tumor angiogenesis, arthritis and atherosclerosis by anti-Flt1. *Nat Med* 2002;8(8):831-40.
42. Gale NW, Yancopoulos GD. Growth factors acting via endothelial cell-specific receptor tyrosine kinases: VEGFs, angiopoietins, and ephrins in vascular development. *Genes Dev* 1999;13(9):1055-66.
- 5 43. Carmeliet P. Mechanisms of angiogenesis and arteriogenesis. *Nat Med* 2000;6(4):389-95.
44. Breiteneder-Geleff S, Soleiman A, Kowalski H, Horvat R, Amann G, Kriehuber E, Diem K, Weninger W, Tschachler E, Alitalo K and others. Angiosarcomas express mixed endothelial phenotypes of blood and lymphatic capillaries: podoplanin as a specific marker for lymphatic endothelium. *Am J Pathol* 1999;154(2):385-94.
- 10 45. Wigle JT, Oliver G. Prox1 function is required for the development of the murine lymphatic system. *Cell* 1999;98(6):769-78.
- 15 46. Jackson DG, Prevo R, Clasper S, Banerji S. LYVE-1, the lymphatic system and tumor lymphangiogenesis. *Trends Immunol* 2001;22(6):317-21.
47. Veikkola T, Jussila L, Makinen T, Karpanen T, Jeltsch M, Petrova TV, Kubo H, Thurston G, McDonald DM, Achen MG and others. Signalling via vascular endothelial growth factor receptor-3 is sufficient for lymphangiogenesis in transgenic mice. *Embo J* 2001;20(6):1223-31.
- 20 48. Sleeman JP, Krishnan J, Kirkin V, Baumann P. Markers for the lymphatic endothelium: in search of the holy grail? *Microsc Res Tech* 2001;55(2):61-9.
49. Witte MH, Bernas MJ, Martin CP, Witte CL. Lymphangiogenesis and lymphangiodysplasia: from molecular to clinical lymphology. *Microsc Res Tech* 2001;55(2):122-45.
- 25 50. Karpanen T, Egeblad M, Karkkainen MJ, Kubo H, Yla-Herttuala S, Jaattela M, Alitalo K. Vascular endothelial growth factor C promotes tumor lymphangiogenesis and intralymphatic tumor growth. *Cancer Res* 2001;61(5):1786-90.
- 30 51. Mandriota SJ, Jussila L, Jeltsch M, Compagni A, Baetens D, Prevo R, Banerji S, Huarte J, Montesano R, Jackson DG and others. Vascular endothelial growth factor-C-mediated lymphangiogenesis promotes tumour metastasis. *Embo J* 2001;20(4):672-82.

52. Skobe M, Hawighorst T, Jackson DG, Prevo R, Janes L, Velasco P, Riccardi L, Alitalo K, Claffey K, Detmar M. Induction of tumor lymphangiogenesis by VEGF-C promotes breast cancer metastasis. *Nat Med* 2001;7(2):192-8.
53. Stacker SA, Caesar C, Baldwin ME, Thornton GE, Williams RA, Prevo R, Jackson DG, Nishikawa S, Kubo H, Achen MG. VEGF-D promotes the metastatic spread of tumor cells via the lymphatics. *Nat Med* 2001;7(2):186-91.
54. Alitalo K, Carmeliet P. Molecular mechanisms of lymphangiogenesis in health and disease. *Cancer Cell* 2002;1(3):219-27.
55. Oliver G, Detmar M. The rediscovery of the lymphatic system: old and new insights into the development and biological function of the lymphatic vasculature. *Genes Dev* 2002;16(7):773-83.
56. Schneider M, Othman-Hassan K, Christ B, Wilting J. Lymphangioblasts in the avian wing bud. *Dev Dyn* 1999;216(4-5):311-9.
57. Kubo H, Cao R, Brakenhielm E, Makinen T, Cao Y, Alitalo K. Blockade of vascular endothelial growth factor receptor-3 signaling inhibits fibroblast growth factor-2-induced lymphangiogenesis in mouse cornea. *Proc Natl Acad Sci U S A* 2002;99(13):8868-73.
58. Folkman J. A mechanism for regulation of lymphangiogenesis independent of angiogenesis. Bethesda; 2002.
59. Baldwin M, Catimel B, Nice EC, Roufail S, Hall NE, Stenvers KL, Karkkainen MJ, Alitalo K, Stacker SA, M.G. A. The Specificity of Receptor Binding by Vascular Endothelial Growth Factor-D Is Different in Mouse and Man. *J. Biol. Chem.* 2001;276(22):19166-19171.
60. Boardman KC, Swartz MA. Lymphangiogenesis in a mouse tail model. 2001. p 671-672.
61. Ikomi F, Ohhashi T. Immunohistochemical expression of VEGF receptors on new vascularization of collecting lymphatics in mice. In: Editore M, editor; 2001; Sidney, Australia. p 387-390.
62. Reynolds BA, Weiss S. Generation of neurons and astrocytes from isolated cells of the adult mammalian central nervous system. *Science* 1992;255(5052):1707-10.
63. Gage FH. Mammalian neural stem cells. *Science* 2000;287(5457):1433-8.
64. Yamamoto S, Yamamoto N, Kitamura T, Nakamura K, Nakafuku M. Proliferation of parenchymal neural progenitors in response to injury in the adult rat spinal cord. *Exp Neurol* 2001;172(1):115-27.

65. Cao Q, Benton RL, Whittemore SR. Stem cell repair of central nervous system injury. *J Neurosci Res* 2002;68(5):501-10.
66. Arvidsson A, Collin T, Kirik D, Kokaia Z, Lindvall O. Neuronal replacement from endogenous precursors in the adult brain after stroke. *Nat Med* 2002;8(9):963-70.
- 5 67. Kempermann G, Kuhn HG, Gage FH. Genetic influence on neurogenesis in the dentate gyrus of adult mice. *Proc Natl Acad Sci U S A* 1997;94(19):10409-14.
68. Kuhn HG, Palmer TD, Fuchs E. Adult neurogenesis: a compensatory mechanism for neuronal damage. *Eur Arch Psychiatry Clin Neurosci* 2001;251(4):152-8.
- 10 69. Gritti A, Parati EA, Cova L, Frolichsthal P, Galli R, Wanke E, Faravelli L, Morassutti DJ, Roisen F, Nickel DD and others. Multipotential stem cells from the adult mouse brain proliferate and self-renew in response to basic fibroblast growth factor. *J Neurosci* 1996;16(3):1091-100.
70. Takagi Y, Mitsui A, Nishiyama A, Nozaki K, Sono H, Gon Y, Hashimoto N, Yodoi J. Overexpression of thioredoxin in transgenic mice attenuates focal ischemic brain damage. *Proc Natl Acad Sci U S A* 1999;96(7):4131-6.
- 15 71. Tamaki S, Eckert K, He D, Sutton R, Doshe M, Jain G, Tushinski R, Reitsma M, Harris B, Tsukamoto A and others. Engraftment of sorted/expanded human central nervous system stem cells from fetal brain. *J Neurosci Res* 2002;69(6):976-86.
- 20 72. Wu S, Suzuki Y, Kitada M, Kataoka K, Kitaura M, Chou H, Nishimura Y, Ide C. New method for transplantation of neurosphere cells into injured spinal cord through cerebrospinal fluid in rat. *Neurosci Lett* 2002;318(2):81-4.
73. Lang-Lazdunski L, Matsushita K, Hirt L, Waeber C, Vonsattel JP, Moskowitz MA, Dietrich WD. Spinal cord ischemia. Development of a model in the mouse. *Stroke* 2000;31(1):208-13.
- 25 74. Rapalino O, Lazarov-Spiegler O, Agranov E, Velan GJ, Yoles E, Fraidakis M, Solomon A, Gepstein R, Katz A, Belkin M and others. Implantation of stimulated homologous macrophages results in partial recovery of paraplegic rats. *Nat Med* 1998;4(7):814-21.